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SUBSTITUTE FORM PTO-1449 (MODIFIED)		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE		Attorney Docket No.	50206-013003
				Serial No.	10/654,796
				Applicant	Nicholas P. Barker et al.
				Filing Date	September 3, 2003
				Group	1614
				IDS Filed	June 23, 2004
				Customer No.	21559

U.S. PATENTS						
Examiner's Initials	Patent Number	Issue Date	Patentee	Class	Subclass	Filing Date (If Appropriate)
RPJ	4,647,454	03/03/87	Cymbalista			
	5,349,001	09/20/94	Greenwald et al.			
	5,359,030	10/25/94	Ekwuribe			
	5,382,657	01/17/95	Karasiewicz et al.			
	5,446,090	08/29/95	Haris			
↓	6,296,844	10/02/01	Takahashi			

FOREIGN PATENT OR PUBLISHED FOREIGN PATENT APPLICATION						
Examiner's Initials	Document Number	Publication Date	Country or Patent Office	Class	Subclass	Translation (Yes/No)
BDJ	WO 98/16255 A2	04/23/98	PCT			
↓	WO 00/66137 A1	11/9/00	PCT			
↓	WO 02/32414 A2	04/25/02	PCT			

OTHER DOCUMENTS (INCLUDING AUTHOR, TITLE, DATE, PLACE OF PUBLICATION)						
BDJ	Bailon et al., "Rational design of a potent, long-lasting form of interferon: A 40 kDa branched polyethylene glycol-conjugated interferon α 2a for the treatment of hepatitis C," <i>Bioconjugate Chem.</i> 12:195-202 (2001).					
	Burgess et al., "Abnormal surface distribution of the human asialoglycoprotein receptor in cirrhosis," <i>Hepatology</i> 15:702-706 (1992).					
	Cutrone et al., "Identification of critical residues in bovine IFNAR-1 responsible for interferon binding," <i>J. Biol. Chem.</i> 276:17140-17148 (2001):					
↓	Dotzauer et al., "Hepatitis A virus-specific immunoglobulin A mediates infection of hepatocytes with hepatitis A virus via the asialoglycoprotein receptor," <i>J. Virology</i> 74:10950-10957 (2000).					
EXAMINER	<i>Bruce J. Harris</i>			DATE CONSIDERED	2/26/06	
EXAMINER: Initial citation considered. Draw line through citation if not in conformance and not considered. Include copy of this form with the next communication to applicant.						



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		INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Use several sheets if necessary)		Serial No.	10/654,796
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<p>BDH</p> <p>Eisenberg et al., "Asialoglycoprotein receptor in human isolated hepatocytes from normal liver and its apparent increase in liver with histological alterations," <i>J. Hepatol.</i> 13:305-309 (1991).</p>					
<p>Eto et al., "Enhanced inhibition of hepatitis B virus production by asialoglycoprotein receptor-directed interferon," <i>Nature Medicine</i> 5:577-581 (1999).</p>					
<p>Glue et al. "Pegylated interferon- α 2b: pharmacokinetics, pharmacodynamics, safety, and preliminary efficacy data," <i>Clinical Pharmacology & Therapeutics</i> 68:556-567 (2000).</p>					
<p>Grace et al., "Structural and biologic characterization of pegylated recombinant IFN- α 2b," <i>J Interferon Cytokine Res.</i> 21:1103-1115 (2001).</p>					
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<p>Karpusas et al., "The crystal structure of human interferon β at 2.2-A resolution," <i>Proc. Natl. Acad. Sci. USA</i> 94:11813-11818 (1997).</p>					
<p>Kasama et al., "Pharmacokinetics and biologic activities of human native and asialointerferon- βs," <i>J. Interferon Cytokine Res.</i> 15:407-415 (1995).</p>					
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<p>Mizuno et al., "Distribution of asialoglycoprotein receptor in human hepatocellular carcinoma," <i>Liver</i> 13:80-85 (1993).</p>					
<p>Monkarsh et al., "Positional isomers of monopegylated interferon α-2a: Isolation, characterization, and biological activity," <i>Anal. Biochem.</i> 247:434-440 (1997).</p>					
<p>Nyman et al., "Structural characterisation of N-linked and O-linked oligosaccharides derived from interferon- α 2b and interferon- α 14c produced by Sendai-virus-induced human peripheral blood leukocytes," <i>Eur. J. Biochem.</i> 253:485-493 (1998).</p>					
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OTHER DOCUMENTS (INCLUDING AUTHOR, TITLE, DATE, PLACE OF PUBLICATION)					
<p><i>BDH</i></p> <p>Pepinsky et al., "Improved pharmacokinetic properties of a polyethylene glycol-modified form of interferon-β-1a with preserved in vitro bioactivity," <i>J. Pharmacol. Exp. Ther.</i> 297:1059-1066 (2001).</p>					
<p>Qin et al., "Interferon-β gene therapy inhibits tumor formation and causes regression of established tumors in immune-deficient mice," <i>Proc. Natl. Acad. Sci. USA</i> 95:14411-14416 (1998).</p>					
<p>Qin et al., "Human and mouse IFN-β gene therapy exhibits different anti-tumor mechanisms in mouse models," <i>Mol. Therapy</i> 4:356-364 (2001).</p>					
<p>Schering-Plough Research Institute, "Toxicologist's Review of PEGylated Interferon-α 2b (PEG-IFN, PEG-INTRON)," December 19, 2000.</p>					
<p>Tada et al., "Systemic IFN-β gene therapy results in long-term survival in mice with established colorectal liver metastases," <i>J. Clin. Invest.</i> 108:83-95 (2001).</p>					
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EXAMINER <i>Buried 9/11/05</i>		DATE CONSIDERED <i>2/26/06</i>			
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